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The Prevalence and Correlates of Eating Disorders in the National Comorbidity Survey Replication

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Abstract

Background— Little population-based data exist on the prevalence or correlates of eating disorders.

Methods— Prevalence and correlates of eating disorders from the National Comorbidity Replication, a nationally representative face-to-face household survey (n_9282), conducted in 2001–2003, were assessed using the WHO Composite International Diagnostic Interview.

Results— Lifetime prevalence estimates of DSM-IV anorexia nervosa, bulimia nervosa, and binge eating disorder are .9%, 1.5%, and 3.5% among women, and .3% .5%, and 2.0% among men. Survival analysis based on retrospective age-of-onset reports suggests that risk of bulimia nervosa and binge eating disorder increased with successive birth cohorts. All 3 disorders are significantly comorbid with many other DSM-IV disorders. Lifetime anorexia nervosa is significantly associated with low current weight (body-mass index18.5), whereas lifetime binge eating disorder is associated with current severe obesity (body-mass index < _40). Although most respondents with 12-month bulimia nervosa and binge eating disorder report some role impairment (data unavailable for anorexia nervosa since no respondents met criteria for 12-month prevalence), only a minority of cases ever sought treatment.

Conclusions— Eating disorders, although relatively uncommon, represent a public health concern because they are frequently associated with other psychopathology and role impairment, and are frequently under-treated.

Keywords

Anorexia nervosa; binge eating disorder; bulimia nervosa; eating disorders; epidemiology; national comorbidity survey replication (NCS-R)

Two eating disorders—anorexia nervosa and bulimia nervosa—are recognized as diagnostic entities in the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) (American Psychiatric Association 1994); a third category, binge eating disorder, is proposed in DSM-IV as a possible new diagnostic entity. However, data are incomplete on the prevalence of these 3 disorders in the general population. The prevalence of anorexia nervosa has been investigated mainly in samples of young women in Europe and North America, where the average point prevalence has been .3% (Hoek and van Hoeken 2003; Favaro et al 2004). The lifetime prevalence among adult women has been reported as .5%-.6% in 2 large population-based surveys in the United States (Walters and Kendler 1995) and Canada (Garfinkel et al 1996); the latter study found a prevalence of anorexia nervosa among adult men of .1%. The lifetime prevalence of bulimia nervosa in adult women has been estimated as 1.1%-2.8% in 3 large population based surveys in New Zealand (Bushnell et al 1990), the United States (Kendler et al 1991), and Canada (Garfinkel et al 1995). For men, the lifetime prevalence of bulimia nervosa was estimated at .1% in the Canadian study and .2% in the New Zealand study, but the point prevalence of bulimia nervosa in a study in Austria was reported as .5% (Kinzl et al 1999b). For the case of binge eating disorder, 2 population-based telephone interview surveys of adults in Austria estimated the point prevalence as 3.3% among women (Kinzl et al 1999a) and .8% among men (Kinzl et al 1999b). Other studies of binge eating

disorder have been limited to specific populations (e.g., young women) or were based only on questionnaires, rather than personal interviews (Streigel-Moore and Franko 2003;Favaro et al 2004).

Population-based interview data are needed to ascertain the prevalence of the 3 eating disorders as well as to provide data on age-of-onset distributions, duration, and association with sociodemographics and body-mass index (BMI). Population data could also address the question of cohort effects—whether the incidence of eating disorders has changed in recent decades. Also of interest is the association of eating disorders with other mental disorders, with measures of disability, and with history of mental health treatment. Finally, population-based data may be useful in examining alternative definitions of eating disorder syndromes in order to determine which definitions are most meaningful as markers of psychopathology. To address these questions, we analyzed data from the recently completed National Comorbidity Survey Replication (NCS-R).

Methods and Materials

Sample

The NCS-R is a nationally representative survey of the US household population that was administered face-to-face to a sample of 9282 English-speaking adults ages 18 and older between February 2001 and December 2003 (Kessler and Merikangas 2004). The response rate was 70.9%. The sample was based on a multi-stage clustered area probability design. Recruitment featured an advance letter and Study Fact Brochure followed by in-person interviewer visits to obtain informed consent. Consent was verbal rather than written in order to parallel the consent procedures in the baseline NCS (Kessler et al 1994). Respondents were given a \$50 financial incentive for participation. The Human Subjects Committees of both Harvard Medical School and the University of Michigan approved these recruitment and consent procedures.

The survey was administered in 2 parts. Part I included the core diagnostic assessment and was administered to all respondents. Part II assessed additional disorders and correlates of disorders. Part II was administered to a subset of 5692 respondents consisting of all those who met lifetime criteria for a Part I disorder plus a probability sample of other respondents. Disorders of secondary interest were administered to probability sub-samples of the Part II sample. Eating disorders were among the latter disorders.

The analyses reported here were carried out in a sub-sample of 2980 Part II respondents who were randomly assigned to have an assessment of eating disorders. Data records in this subsample were weighted to adjust for the over-sampling of Part I respondents with a mental disorder, differential probabilities of selection within households, systematic non-response, and residual socio-demographic-geographic differences between the sample and the 2000 Census. NCS-R sampling and weighting are discussed in more detail elsewhere (Kessler et al 2004b).

Diagnostic Assessment

NCS-R diagnoses were based on Version 3.0 of the World Health Organization Composite International Diagnostic Interview (CIDI) (Kessler and Ustun 2004), a fully structured layadministered diagnostic interview that generates diagnoses according to both ICD-10 and DSM-IV criteria. DSM-IV criteria were used in the current report. Core disorders included the three broad classes of disorder assessed in previous CIDI surveys (anxiety disorders, mood disorders, and substance disorders) plus a group of disorders that share a common feature of difficulties with impulse control (e.g., intermittent explosive disorder, attention-deficit/

hyperactivity disorder, retrospectively reported childhood oppositional-defiant disorder, and conduct disorder). Diagnostic hierarchy rules and organic exclusion rules were used in making all diagnoses. As detailed elsewhere (Kessler et al 2004a,2005), good concordance was found between these core CIDI diagnoses and diagnoses based on the Structured Clinical Interview for DSM-IV (SCID) (First et al 2002) in a probability sub-sample of NCS-R respondents who were administered clinical reappraisal interviews. The area under the receiver operator characteristic curve was in the range of .65–.81 for anxiety disorders, .75 for major depressive episode, .62–.88 for substance disorders, and .76 for any anxiety, mood, or substance disorder. No clinical reappraisal interviews were carried out for the impulse-control disorders, as these were not core NCS-R disorders.

For the present study, questions from the CIDI were used to assign diagnoses of anorexia nervosa, bulimia nervosa, and binge eating disorder based on DSM-IV criteria. The full diagnostic algorithms for all 3 disorders, together with a sensitivity analysis using alternative, narrower definitions of bulimia nervosa and binge eating disorder, are presented as supplemental material available online with the electronic version of this article and at www.hcp.med.harvard.edu/ncs//eating.php; the corresponding CIDI questions used to operationalize the criteria are available at www.hcp.med.harvard.edu/ncs.

Most of the CIDI questions closely paralleled the DSM-IV criteria, but to meet criteria for binge eating disorder, DSM-IV requires a minimum of 6 months of regular eating binges, whereas the CIDI asked only whether the individual experienced 3 months of symptoms. Thus, individuals displaying more than 3 months, but less than 6 months, of regular binge eating would be classified as having binge eating disorder in our algorithm, but not in DSM-IV. Also of note is that for binge eating episodes in bulimia nervosa and binge eating disorder, DSM-IV requires assessment of loss of control, and for binge eating disorder requires marked distress regarding binge eating; these items were assessed in the CIDI by a series of questions about attitudes and behaviors that are indicators of loss of control and of distress, rather than by direct questions.

In addition to the 3 eating disorders, we also defined 2 provisional entities. The first was "subthreshold binge eating disorder," defined as a) binge eating episodes, b) occurring at least twice a week for at least 3 months, and c) not occurring solely during the course of anorexia nervosa, bulimia nervosa, or binge eating disorder. Thus, subthreshold binge eating disorder did not require DSM-IV criterion B (3 of 5 features associated with binge eating) or C (marked distress regarding binge eating for binge eating disorder). The second was "any binge eating," also defined as a) binge eating episodes (again, not requiring DSM-IV criteria B and C), b) occurring at least twice a week for at least 3 months, but c) lacking the hierarchical exclusion criterion if the individual simultaneously exhibited another eating disorder. In other words, any binge eating was diagnosed regardless of whether or not the individual simultaneously met criteria for any of the other 3 eating disorders or for subthreshold binge eating disorder. This entity thus included all cases of bulimia nervosa, binge eating disorder, and subthreshold binge eating disorder, as well as cases of anorexia nervosa with binge eating. Full diagnostic algorithms for these 2 provisional entities, together with a sensitivity analysis parallel to that above, are presented as supplemental material available with the online version of this article and at www.hcp.med.harvard.edu/ncs//eating.php.

In summary, we examined a total of 5 conditions—2 official DSM-IV disorders (anorexia nervosa and bulimia nervosa), 1 proposed DSM-IV disorder (binge eating disorder), and 2 provisional entities that partially overlapped with 1 or more of the previous 3 disorders. Although in the following text we refer to these 5 conditions collectively as "disorders" for simplicity, the reader should bear in mind that they vary in terms of their level of general acceptance.

As indicated above, our criteria allowed that individuals could display more than one lifetime diagnosis of an eating disorder. We used data from the CIDI regarding time of onset and recency (i.e., the time when the disorder was last present) to apply diagnostic hierarchies, so that bulimia nervosa, binge eating disorder, and subthreshold binge eating disorder were not diagnosed in the presence of anorexia nervosa; and so that binge eating disorder and subthreshold binge eating disorder were not diagnosed in the presence of bulimia nervosa. Because the CIDI provides information only about onset and recency of a disorder, individuals with an episode of a given eating disorder occurring only in between two or more discrete episodes of a hierarchically exclusionary disorder (e.g., anorexia nervosa) would not have been diagnosed with that disorder.

For individuals meeting criteria for any of the 5 five disorders, the CIDI assessed age of onset, recency, years with the disorder, and professional help-seeking. Respondents with 12-month prevalence (that is, individuals who met criteria for the eating disorder at any time within the 12 months before interview) were additionally administered the Sheehan Disability Scales (Leon et al 1997) to assess the severity of recent episodes and were asked about treatment in the past 12 months.

Statistical Analyses

Cross-tabulations were used to estimate prevalence, disability, and treatment. The actuarial method (Wolter 1985) was used to estimate age-of-onset curves. Discrete-time survival analysis with the person-year as the unit of analysis (Willett and Singer 1993) using logistic regression (Hosmer and Lemeshow 2000) was used to estimate cohort effects. Logistic regression was also used to study socio-demographic correlates and comorbidity. Logits and their 95% confidence intervals were converted into odd ratios by exponentiation for ease of interpretation. Standard errors and significance tests were estimated using the Taylor series linearization method (Wolter 1985) implemented in the SUDAAN software system (Research Triangle Institute 2002) to adjust for the weighting and clustering of the NCS-R data. Multivariate significance of predictor sets was evaluated using Wald _ 2 tests based on design-corrected coefficient variance-covariance matrices. Statistical significance was evaluated using 2-tailed .05-level tests; it should be noted that this level, which was pre-specified for all NCS-R analyses, does not correct for multiple comparisons and thus underestimates the overall type I error rate.

Results Prevalence

Lifetime prevalence estimates of anorexia nervosa, bulimia nervosa, binge eating disorder, subthreshold binge eating disorder, and any binge eating were .6%, 1.0%, 2.8%, 1.2%, and 4.5% (Table 1). Lifetime prevalence was consistently $1\frac{3}{4}$ to 3 times as high among women as men for the 3 eating disorders (z_2.2–2.8, P_.029–.005), 3 times as high among men as women for subthreshold binge eating disorder (z_3.3, P_.001), and approximately equal among women and men for any binge eating (z_1.2., P_.219). No 12-month cases of anorexia nervosa were found in the sample. The 12-month prevalence estimates of the other 4 disorders were considerably lower than the lifetime estimates, although with similar sex ratios. Estimates of cumulative lifetime risk by age 80, based on retrospective age-of-onset reports (Figure 1), were 0.6% for anorexia nervosa, 1.1% for bulimia nervosa, 3.9% for binge eating disorder, 1.4% for subthreshold binge eating disorder, and 5.7% for any binge eating.

Age of Onset and Persistence

Median age of onset of the five disorders ranged from 18–21 years (Table 2). The period of onset risk was shorter for anorexia nervosa than for the other disorders, with the earliest cases of the other disorders beginning about 5 years earlier than those of anorexia nervosa (ages 10

vs. 15), and no cases of anorexia nervosa beginning after the mid-20s—whereas some cases of the other disorders began at a much older age (Figures 1 and 2).

The mean number of years with anorexia nervosa (1.7 years) was significantly lower than for either bulimia nervosa $(8.3; t_4.1, P_.001)$, binge eating disorder $(8.1; t_2.9, P_.006)$, subthreshold binge eating disorder $(7.2; t_2.6, P_.013)$, or any binge eating $(8.7; t_2.9, P_.005)$ (Table 2). Consistent with these differences in duration, 12-month persistence, defined as 12-month prevalence among lifetime cases, was lowest for anorexia nervosa (.0%) and higher for bulimia nervosa (30.6%), binge eating disorder (44.2%), subthreshold binge eating (47.2%), and any binge eating (46.9%).

Cohort Effects

Consistent inverse associations between cohort (age at interview) and lifetime risk were found in survival analyses of all 5 disorders (Table 3). However, the odds ratios in younger (ages 18–29, 30–44) versus older (60_) cohorts were significantly higher for all comparisons only for bulimia nervosa, binge eating disorder, and any binge eating.

Association with Body-Mass Index

Individuals with a lifetime diagnosis of anorexia nervosa displayed a significantly lower current BMI—with a greater prevalence of a current BMI of $_$ 18.5, and a lower prevalence of a current BMI $_$ 40—than respondents without any eating disorder (Table 4). The reverse pattern was found for binge eating disorder, with a significantly higher prevalence of BMI of $_$ 40 among individuals with binge eating disorder than respondents without any eating disorder. Any binge eating was also associated with severe obesity, but this finding was attributable entirely to cases of binge eating disorder.

Twelve-Month Role Impairment

Role impairment was assessed only for 12-month cases; since there were no 12-month cases of anorexia nervosa, our analysis was limited to the other 4 disorders. The majority of respondents with bulimia nervosa, binge eating disorder, or any binge eating reported at least some role impairment (mild, moderate, or severe) in at least 1 role domain (53.1%–78.0%), but only 21.8% of respondents with subthreshold binge eating disorder reported this degree of impairment (Table 5). Severe role impairment was much less common, and ranged from 3.4% in subthreshold binge eating to 16.3% in bulimia nervosa, with no significant differences in prevalence among groups.

Comorbidity

More than half (56.2%) of respondents with anorexia nervosa, 94.5% with bulimia nervosa, 78.9% with binge eating disorder, 63.6% with subthreshold binge eating disorder, and 76.5% with any binge eating met criteria for at least 1 of the core DSM-IV disorders assessed in the NCS-R (Table 6). Eating disorders were positively related to almost all of the core DSM-IV mood, anxiety, impulse-control, and substance use disorders after controlling for age, sex, and race-ethnicity, with 89% of the odds ratios for the association between individual eating disorders and individual comorbid conditions greater than 1.0 and 67% significant at the .05 level. The odds ratios were consistently largest, though, for bulimia nervosa, with a median (and inter-quartile range in parentheses) odds ratio of 4.7 (4.3–7.5), next highest for binge eating disorder (3.2 [2.6–3.7]) and any binge eating (3.2 [2.4–3.8]), and smaller for anorexia nervosa (2.1 [1.2–2.9]) and subthreshold binge eating disorder (2.2 [1.1–2.9]). No single class of disorders stood out as showing consistently or markedly higher comorbidity with eating disorders.

Treatment

A majority of respondents with anorexia nervosa, bulimia nervosa, and binge eating disorder (50.0%–63.2%) received treatment for emotional problems at some time in their lives, with the most common site of treatment being the general medical sector for anorexia nervosa (45.3%) and binge eating disorder (36.3%), and the mental health specialty sector for bulimia nervosa (48.2% for psychiatrist and 48.3% for other mental health) (Table 7). However, smaller proportions sought treatment specifically for their bulimia nervosa (43.2%) or binge eating disorder (43.6%). Only 15.6% of respondents with 12-month bulimia nervosa and 28.5% with 12-month binge eating disorder received treatment for emotional problems in the 12 months before interview, with the most common site of treatment being the general medical sector, and similar proportions received 12-month treatment specifically for their bulimia nervosa or binge eating disorder.

Supplemental data are available with the electronic version of this article and online at www.hcp.med.harvard.edu/ncs/eating.php.

Discussion

In a population-based survey of American households—the first nationally representative study of eating disorders in the United States—we found estimates of lifetime prevalence for eating disorders that are broadly consistent with earlier data. However, we found a surprisingly high proportion of men with anorexia nervosa and bulimia nervosa (representing approximately one-fourth of cases of each of these disorders). By contrast, clinical and case registry studies (Fairburn and Beglin 1990;Hoek and van Hoeken 2003) report that fewer than 10% men among cases of these disorders, and population-based studies report a 15% proportion of men for anorexia nervosa (Garfinkel et al 1996) and 8%–10% of men for bulimia nervosa (Bushnell et al 1990;Garfinkel et al 1995). Note, however, that estimates from population-based studies, including ours, are unstable because they involve small numbers of men with eating disorders (no more than 5 men with either disorder in any study).

Our findings provide unique data regarding the lifetime duration of eating disorders, and the onset and duration of binge eating disorder, together with extensive information on sociodemographic features of individuals with all 5 disorders. Also, our study provides support for the common impression that the incidence of bulimia nervosa has increased significantly in the second half of the twentieth century (Kendler et al 1991;Hoek and van Hoek 2003), and it provides the first data showing a similar trend for binge eating disorder. Nevertheless, there are some data suggesting that the incidence of bulimia nervosa may be leveling off in recent years (Currin et al 2005). Whether the incidence of anorexia nervosa has increased over time is unclear and subject to debate. We failed to find a significant increase, but had little power to detect such a trend; case registry study data have yielded conflicting findings and interpretations (Fombonne, 1995;Lucas et al 1999;Hoek and van Hoeken 2003;Currin et al 2005).

We found that lifetime anorexia nervosa is associated with a low current BMI, a finding consistent with follow-up studies of clinical samples of individuals with anorexia nervosa showing that low weight often persists after resolution of the disorder (Steinhausen 2002). By contrast, binge eating disorder was found to be strongly associated with current severe obesity (BMI_40)—a finding also consistent with earlier reports (de Zwaan 2001;Streigel-Moore and Franko 2003;Hudson et al 2006). Although the causal pathways responsible for this latter association are unclear, shared familial factors (such as shared genes or shared family environmental exposures) are likely at least partly responsible (Hudson et al 2006).

We also assessed role impairment in all disorders except anorexia nervosa, where analysis was precluded because no 12-month cases were identified. While the majority of respondents with bulimia nervosa, binge eating disorder, or any binge eating reported at least some role impairment in at least 1 role domain, only 21.8% of respondents with subthreshold binge eating disorder reported any role impairment. Severe role impairment was uncommon in all conditions. It is important to note, though, that participants may possibly have under-reported role impairment due to factors such as minimization, shame, secrecy, or lack of insight stemming from the ego-syntonicity of symptoms.

Less than half of individuals with bulimia nervosa or binge eating disorder had ever sought treatment for their eating disorder (a measure not assessed for anorexia nervosa), although the majority of individuals with all 3 disorders had received treatment at some point for some emotional problem. This finding, coupled with the observation that physicians infrequently assess patients for binge eating (Crow et al 2004) and often fail to recognize bulimia nervosa and binge eating disorder (Johnson et al 2001), highlights the importance of querying patients about eating problems even when they do not include such problems among their presenting complaints.

We found a high prevalence of lifetime comorbid psychiatric disorders in individuals with all disorders except subthreshold binge eating disorder, although this finding was less pronounced for anorexia nervosa. These results are again generally consistent with those reported in previous population-based studies for anorexia nervosa (Garfinkel et al 1996), bulimia nervosa (Kendler et al 1991;Bushnell et al 1994;Garfinkel et al 1995;Rowe et al 2002), binge eating behavior (Vollrath et al 1992;Angst 1988;Bulik et al 2002), and regular binge eating without compensatory behaviors (Reichborn-Kjennerud et al 2004b), as well as in previous studies of clinical populations for anorexia nervosa, bulimia nervosa, and binge eating disorder (Hudson et al 1987;Halmi et al 1991;Johnson et al 2001;Godart et al 2002;Kaye et al 2004;McElroy et al 2005). The cause for the high levels of comorbidity is not known, although there is evidence that the co-occurrence of eating disorders with mood disorders may be caused in part by common familial (Mangweth et al 2003) or genetic factors (Walters et al 1992;Wade et al 2000).

Several findings in this study are particularly noteworthy. First, we found that anorexia nervosa displayed a significantly shorter lifetime duration and lower 12-month persistence, as well as lower overall levels of comorbidity, than either bulimia nervosa or binge eating disorder. These findings contrast with previous studies (Steinhausen 2002) that have conceptualized anorexia nervosa as a chronic and malignant condition. This discrepancy may be due to the fact that our population-based method identified individuals with milder cases of anorexia nervosa who might have been missed in previous follow-up studies, which were based largely on clinical samples. Alternatively, our population-based method might have missed more severe cases of anorexia nervosa, either because they were unavailable, unreachable, hospitalized, or unwilling to participate in an interview about emotional problems. Parenthetically, we would note that while we found no cases of current anorexia nervosa in our study, 15.6% of the individuals with a lifetime diagnosis of anorexia nervosa still had a current BMI of less than 18.5 at the time of interview. Indeed, these individuals (3 cases) were all below 85% of ideal body weight, thus meeting our operationalization for DSM-IV criterion A for anorexia nervosa. However, all of these individuals failed to meet at least one of the other criteria for anorexia nervosa currently—although our data did not permit an analysis of which specific criteria were lacking in individual cases. Nevertheless, these data suggest that a minority of individuals with past anorexia nervosa may continue to maintain an abnormally low body weight, even though they no longer meet full criteria for anorexia nervosa.

Our findings also provide further evidence for the clinical and public health importance of binge eating disorder. In contrast to some earlier studies suggesting that binge eating disorder might be a relatively transient condition (Cachelin et al 1999;Fairburn et al 2000), the present findings, together with those from another recent study (Pope et al, in press), suggest that this disorder is at least as chronic and stable as anorexia nervosa or bulimia nervosa. Binge eating disorder also appears more common than either of the other two eating disorders, exhibits substantial comorbidity with other psychiatric disorders, and is strongly associated with severe obesity. Collectively, these findings suggest that binge eating disorder represents a public health problem at least equal to that of the other 2 better-established eating disorders, adding support to the case for elevating binge eating disorder from a provisional entity to an official diagnosis in DSM-V.

Subthreshold binge eating disorder, by contrast, was found to be associated with such low impairment and comorbidity that it likely does not merit consideration for inclusion as a DSM disorder. It should be recalled, in this connection, that the main difference between subthreshold binge eating disorder and binge eating disorder is that the former lacks the criterion of distress (see Appendix Table 1 in Supplement 1). These findings suggest that the criterion of distress may be important for defining clinically meaningful forms of binge eating.

Note that subthreshold binge eating disorder may be defined in different ways. For example, relaxing the frequency criteria to less than the average of 2 days per week for 6 months required by DSM-IV identifies groups with characteristics similar to the full disorder (Striegel-Moore et al 2000; Crow et al 2002). We were unable, however, to evaluate these definitions due the nature of the CIDI questions, and instead defined subthreshold binge eating disorder by relaxing criteria other than frequency of binges. Thus, while our definition of subthreshold binge eating disorder does not appear to identify a clinically meaningful entity, other definitions may well do so.

Unlike subthreshold binge eating disorder, the entity "any binge eating" is associated with severe obesity, modest levels of impairment, and high levels of comorbidity with other mental disorders. These features appear to be accounted for cases of bulimia nervosa or binge eating disorder within the "any binge eating" group, given that such features are not shared by those with subthreshold binge eating disorder, and individuals with anorexia nervosa contribute only a small number of cases. The findings for any binge eating are interesting to consider in the light of findings from twin studies of binge eating. These studies have suggested that there are genetic influences on binge eating (Bulik et al 1998) and on binge eating without compensatory behaviors (Reichborn-Kjennerud et al 2004a). On the basis of our findings here, it is tempting to speculate that the heritability of binge eating behavior may be attributable primarily to cases of bulimia nervosa and binge eating disorder—both of which have been shown to be familial (Strober et al 2000; Hudson et al 2006)—rather than to cases of subthreshold binge eating disorder within the group.

Several limitations of the study should be considered. First, some CIDI questions did not precisely mirror the DSM-IV criteria for the various eating disorders, as illustrated by in the diagnostic algorithms discussed in our methods section. Perhaps the most important inconsistency is that, in order to have parallel duration requirements for bulimia nervosa and for binge eating disorder, we required only 3 months of illness for a diagnosis of binge eating disorder, in contrast to the 6 months required by DSM-IV. Thus, it is possible that we may have overestimated the prevalence of binge eating disorder by including some cases with a duration of only 3 to 5 months.

Second, diagnoses were based on unvalidated, fully structured lay interviews where lifetime information was assessed retrospectively. These may be important considerations, given that

an earlier version of the CIDI was found to underdiagnose eating disorders (Thornton et al 1998), possibly because some individuals minimized or denied symptoms. Version 3.0 of the CIDI was designed to reduce this sort of under-reporting by using a number of techniques developed by survey methodologists to reduce embarrassment and other psychological barriers to reporting (Kessler and Ustun 2004)—but these changes necessitated indirect assessments of loss of control and distress, as noted above. In any event, pending validation studies, it would seem prudent to think of the NCS-R estimates as lower bounds on the true prevalence of eating disorders.

Third, in our analyses of the associations between eating disorders and body weight, we possessed only current BMI, rather than maximum or minimum adult BMI, or BMI at the time of the disorder. Thus, we likely underestimated the magnitude of these associations.

Fourth, because recall of earlier experiences may diminish with age, our retrospective assessments may have overestimated the magnitude of cohort effects (Giuffra and Risch 1994). Since cohort effects and age effects are confounded, and no prospective studies have been performed over the period under study, it is not possible to assess the magnitude of this potential bias. Prospective studies will be useful to track possible cohort effects in the future.

Fifth, our results are based on the assumption that any exiting from the population available for sampling was non-informative and that there was no selection bias (in the form of non-response bias) due to sampling from available subjects; these limitations are discussed elsewhere (Hudson et al 2005). For example, the validity of our results would be threatened if the development of eating disorders rendered individuals less likely to be available for sampling, which might occur if there were a high mortality due to eating disorders, or a significant proportion of cases hospitalized at the time of sampling. Although some clinical follow-up studies have suggested substantial mortality for anorexia nervosa (Sullivan 1995;Steinhausen 2002;Keel et al 2003), data from a community case registry study (Iacovino 2004) did not find excess mortality.

Another possible threat to validity would be bias in sampling of available individuals, in that individuals with eating disorders might be more or less likely to participate. However, we carried out a non-response survey to deal with this problem, which offered a larger financial incentive (\$100) to main survey nonrespondents for a short (15-min) telephone interview that assessed diagnostic stem questions. Very little evidence was found that survey respondents and non-respondents differed on stem question endorsement for the NCS-R core anxiety, mood, impulsecontrol, or substance use disorders (Kessler et al 2004b). Thus, it is likely that non-response bias for eating disorders was minimal.

Sixth, while we examined 2 provisional entities in addition to those for which criteria were provided in DSM-IV, we did not examine many other possible entities that lie within the category of Eating Disorder Not Otherwise Specified (Fairburn and Bohn, 2005)—such as subthreshold forms of anorexia nervosa and bulimia nervosa, alternative definitions for subthreshold binge eating disorder (discussed above), purging without either bulimia nervosa or anorexia nervosa (Keel et al 2005), and night eating syndrome (Stunkard et al 2005)—because the questions in the CIDI did not permit evaluation of these conditions.

In conclusion, the lifetime prevalence of the individual eating disorders ranged from 0.6–4.5%; these disorders displayed substantial comorbidity with other DSM-IV disorders and were frequently associated with role impairment. These patterns raise concerns that such a low proportion of individuals with these disorders obtain treatment for their eating problems. As it turns out, though, a high proportion of cases did receive treatment for comorbid conditions. Thus, detection and treatment of eating disorders might be increased substantially if treatment

providers queried patients about possible eating problems, even if the patients did not include such problems among their presenting complaints.

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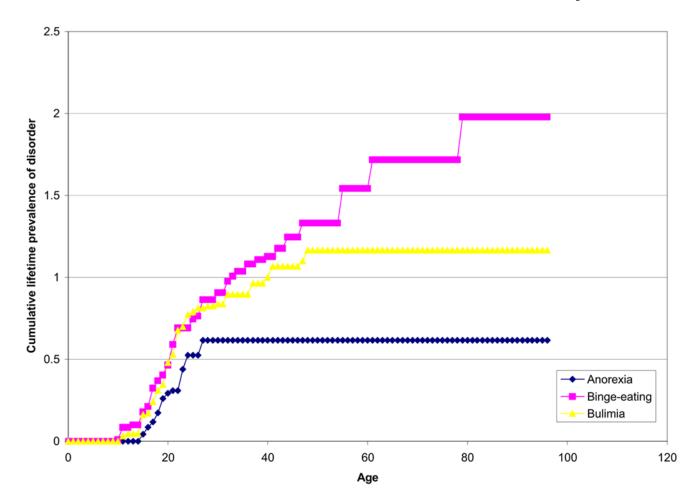


Figure 1. Age-of-onset distributions for DSM-IV eating disorders

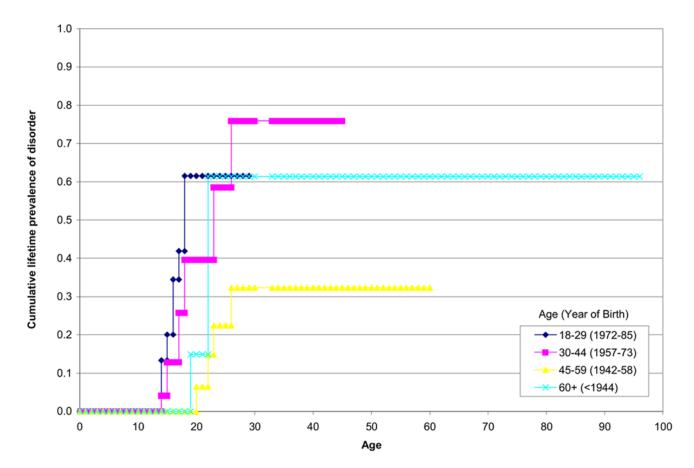


Figure 2. Cohort-specific age-of-onset distributions for DSM-IV Anorexia Nervosa

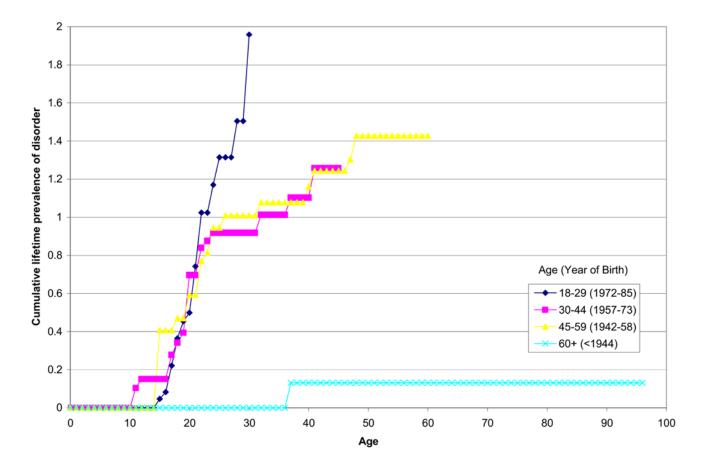


Figure 3. Cohort-specific age-of-onset distributions for DSM-IV Bulimia Nervosa

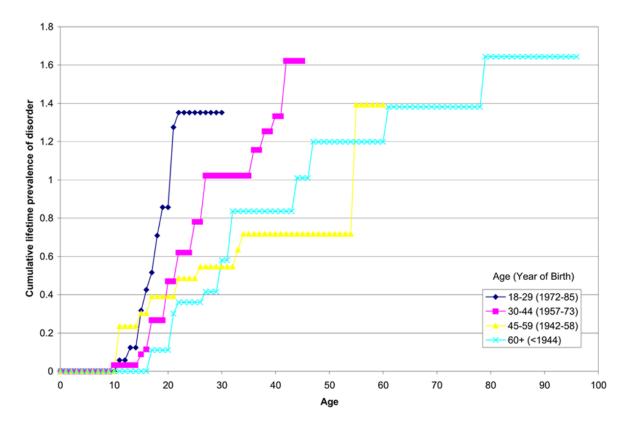


Figure 4. Cohort-specific age-of-onset distributions for DSM-IV Binge-Eating Disorder

Table 1Lifetime and 12-month prevalence estimates of DSM-IV eating disorders and related behavior

	M	ale	Fer	nale	T	otal
	%	(se)	%	(se)	%	(se)
I. Lifetime prevalence						
Anorexia Nervosa	0.3*	(0.1)	0.9*	(0.3)	0.6	(0.2)
Bulimia Nervosa	0.5*	(0.3)	1.5*	(0.3)	1.0	(0.2)
Binge-eating Disorder	2.0*	(0.5)	3.5*	(0.5)	2.8	(0.4)
Subthreshold Binge-eating	1.9*	(0.5)	0.6*	(0.1)	1.2	(0.2)
Any binge-eating behavior II. Twelve-month prevalence	4.0	(0.7)	4.9	(0.6)	4.5	(0.4)
Bulimia Nervosa	0.1*	(0.1)	0.5	(0.2)	0.3	(0.1)
Binge-eating Disorder	0.8*	(0.3)	1.6*	(0.2)	1.2	(0.2)
Subthreshold binge-eating	0.8	(0.3)	0.4	(0.1)	0.6	(0.2)
Any binge-eating behavior	1.7	(0.4)	2.5	(0.3)	2.1	(0.2)
(n)	(12	220)	(17	760)	2	980

^{*} Significant sex difference based on a .05 level, two-sided test.

 $^{{\}cal I}_{\mbox{None}}$ of the respondents met criteria for 12-month Anorexia Nervosa.

	Anorexi	Anorexia Nervosa	Bulimia Nervosa	lervosa	Binge-eating Disorder	g Disorder	Subthreshold Binge-eating	Binge-eating	Any binge	inge-	
	%	(se)	%	(se)	%	(se)	%	(se)	eaung o %	eaung benavior % (se)	(u)
A. Males											
18–29	0.0	:	0.1	0.1	1.4	0.7	2.7	(1.0)	4.1	(1.2)	(288)
30-44	9.0	(0.4)	0.1	0.1	2.5	0.8	2.1	(1.0)	4.6	(1.3)	(403)
45–59	0.0	:	1.3	6.0	2.7	1.2	1.6	(0.8)	4.4	(1.6)	(339)
+09	0.3	(0.3)	0.3	0.3	6.0	0.5	1.1	(0.9)	2.3	(1.0)	(190)
Total	0.3	(0.1)	0.5	0.3	2.0	0.5	1.9	(0.5)	4.0	(0.7)	(1220)
χ^2_3 (age)	2.	2.5^{I}	5.2		3.8	8	1.4		1.9		
B. Females											
18–29	1.1	(0.6)	2.2	0.5	4.2	8.0	0.8	(0.4)	6.2	(0.9)	(417)
30-44	6.0	(0.5)	2.0	9.0	3.7	1.0	8.0	(0.3)	5.9	(1.2)	(564)
45–59	9.0	(0.3)	1.6	0.5	3.4	8.0	0.4	(0.2)	4.4	(0.9)	(462)
+09	8.0	(0.8)	0.0	;	2.4	1.1	0.5	(0.3)	2.9	(1.1)	(317)
Total	6.0	(0.3)			3.5	0.5		(0.1)			(1760)
χ^2_3 (age)	2	2.4	6.9	2	1.1	1	1.3		3.3	3	
C. Total											
18–29	9.0	(0.3)	1.2	0.3	2.9	0.5	1.7	(0.5)	5.2	(0.7)	(705)
30-44	8.0	(0.3)	1.1	0.3	3.1	0.7	1.4	(0.5)	5.3	(0.9)	(296)
45–59	0.3	(0.2)	1.4	0.5	3.1	0.5	6.0	(0.4)	4.4	(0.8)	(801)
+09	9.0	(0.5)	0.1	0.1	1.7	0.7	8.0	(0.4)	2.7	(0.7)	(507)
Total	9.0	(0.2)	1.0	0.2	2.8	0.4	1.2	(0.2)	4.5	(0.4)	2980
χ^2_3 (age)	1	1.8	5.2		2.5	5	2.2		5.2		

* Significant age difference based on a .05 level, χ^2_3 test.

I degrees of freedom = 1

 $\frac{2}{\text{degrees of freedom}} = 2$

NIH-PA Author Manuscript **Table 2** Estimated age of onset and persistence of *DSM-IV* eating disorders and related entities NIH-PA Author Manuscript

	Anorexia Nervosa	Bulimia Nervosa	Binge Eating Disorder	Subthreshold Binge Eating Disorder	Any Binge Eating
Age of onset Mean (se) Median (IQR) Years with episode Mean (se) Median (IQR) 12-month persistence, % (se)	18.9 (0.8) 18.0 (16.0–22.0) 1.7 (0.2) 1.0 (1.0–1.0) 0.0 () (23)	19.7 (1.3) 18.0 (14.0–22.0) 8.3 (1.6) 5.0 (2.0–13.0) 30.6 (7.2) (52)	25.4 (1.2) 21.0 (17.0-32.0) 8.1 (1.1)* 4.0 (1.0-10.0)* 44.2 (6.0)* (115)	22.7 (1.9) 20.0 (17.0–27.0)* 7.2 (2.0)* 2.0 (1.0–10.0)* 47.2 (10.0)* (46)	22.4 (1.1) 20.0 (16.0–27.0)* 8.7 (0.7)* 3.0 (1.0–13.0)* 46.9 (4.1) (192)
	(57)	(35)	(611)	(O+)	

Abbreviations: se, standard error; IQR, interquartile range.

 * Significantly different from a norexia nervosa based on a .05 level, two-sided test.

Appendix table 2
Twelve-month prevalence estimates of DSM-IV eating disorders and related behavior by age and sex

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	Bulimia %	Bulimia Nervosa (se)	Binge-ea %	Binge-eating Disorder % (se)	Subthresh %	Subthreshold Binge-eating % (se)	Any binge-	Any binge-eating behavior % (se)	(u)
A. Males									
18–29	0.0	ı	0.1	(0.1)	9.0	(0.4)	0.7	(0.4)	(288)
30-44	0.0	1	6.0	(0.4)	6.0	(0.4)	1.8	(0.6)	(403)
45–59	0.0	1	1.6	(1.1)	6.0	(0.7)	2.6	(1.3)	(339)
+09	0.3	(0.3)	0.4	(0.4)	0.8	(0.8)	1.5	(0.0)	(190)
Total	0.1	(0.1)	0.8	(0.3)	0.8	(0.3)	1.7	(0.4)	(1220)
χ^2_3 (age)				5.7		6.0		4.9	
B. Females									
18–29	9.0	(0.3)	2.4	(0.7)	9.0	(0.4)	3.6	(0.7)	(417)
30-44	0.7	(0.3)	1.3	(0.4)	0.4	(0.2)	2.5	(9.0)	(564)
45–59	0.7	(0.4)	1.5	(0.0)	0.1	(0.1)	2.3	(0.7)	(462)
+09	0.0	1	1.2	(9.0)	0.3	(0.2)	1.5	(9.0)	(317)
Total	0.5	(0.2)	1.6	(0.2)	0.4	(0.1)	2.5	(0.3)	(1760)
χ^2_3 (age)	1	[.]		3.1		1.8		4.0	
C. Total									
18–29	0.3	(0.2)	1.4	(0.4)	9.0	(0.3)	2.3	(0.4)	(705)
30-44	0.4	(0.2)	1.1	(0.3)	9.0	(0.2)	2.2	(0.4)	(296)
45–59	0.4	(0.2)	1.5	(0.3)	0.5	(0.4)	2.4	(0.5)	(801)
+09	0.1	(0.1)	8.0	(0.4)	0.5	(0.4)	1.5	(0.5)	(207)
Total	0.3	(0.1)	1.2	(0.2)	9.0	(0.2)	2.1	(0.2)	2980
χ^2_3 (age))	6.0		1.6		0.1		1.7	

* Significant age difference based on a .05 level, χ^2 3 test.

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 Table 3

 Inter-cohort differences in lifetime risk of DSM-IV eating disorders and related behavior

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	Anor	Anorexia Nervosa OR (95% CI)	Bulimi OR	Bulimia Nervosa OR (95% CI)	Binge-ea OR	Binge-eating Disorder OR (95% CI)	Subthresh OR	Subthreshold Binge-eating OR (95% CI)	Any binge- OR	Any binge-eating behavior OR (95% CI)
Age (year of birth) 18–29 (1972–	2.0^	(0.7–5.3)	16.8*	(3.0–95.6)	*6.4	(2.1–11.5)	*4.5	(1.8–11.2)	*4.5	(2.6–8.0)
85) 30–44 (1957–	1.8	(0.4–8.2)	*11.8	(1.9–72.6)	3.1	(1.3–7.2)	2.2	(0.8–6.4)	3.0*	(1.7–5.4)
75) 45–59 (1942–	9.0	(0.1–4.5)	13.4*	(2.2–82.2)	2.3*	(1.1–4.9)	1.2	(0.3–4.3)	2.0*	(1.1–3.8)
χ^{28} $60+(<1944)$ χ^{2} $(p-1)$	1.0	2.2 (.104)	1.0	3.8 (.018)*	1.0	6.5 (.001)*	1.0	4.4 (.009)*	1.0	11.3 (.000)*
value)										

* Significant inter-cohort different based on a .05 level, two-sided test, controlling for sex and race-ethnicity.

Collapsing age categories 18–29 and 30–44 (left out category: 35 years or older) results in OR= 2.1 (.9-4.9), χ^2_1 =3.2 (p-value=0.082).

Appendix table 3
Estimated age-of-onset and persistence of DSM-IV eating disorders by lifetime treatment status

	Anorexi	a Nervosa		a Nervosa Means	Binge-eati	ng Disorder
	Mean	(se)	Mean	(se)	Mean	(se)
A. Treated						
Age-of-onset	17.8	(1.1)	24.8	(2.3)	25.1	(2.0)
Years with episode	1.7	(0.3)	8.0*	(1.9)	12.6*	(2.3)
12-month persistence	0.0		44.7	(9.7)	42.1	(7.9)
B. Untreated						
Age-of-onset	19.5	(0.8)	18.1	(1.4)	25.6	(1.6)
Years with episode	1.8	(0.3)	8.6*	(2.1)	4.9*	(0.7)
12-month persistence	0.0		19.8	(7.8)	45.8	(8.9)
C. Total						
Age-of-onset	18.9	(0.8)	19.7	(1.3)	25.4	(1.2)
Years with episode	1.7	(0.2)	8.3*	(1.6)	8.1	(1.1)
12-month persistence	0.0		30.6	(7.2)	44.2	(6.0)
				Iedians		
	Median	(IQR)	Median	(IQR)	Median	(IQR)
A. Treated					***	
Age-of-onset	17.0	(15.0-21.0)	18.0*	(16.0-20.0)	21.0*	(16.0–30.0)
Years with episode	1.0	(1.0-2.0)	4.0*	(2.0-13.0)	5.0*	(2.0-20.0)
12-month persistence	0.0		0.0	(0.0-1.0)	0.0	(0.0-1.0)
B. Untreated						
Age-of-onset	21.0	(17.0-22.0)	17.0*	(13.0-20.0)	19.0*	(16.0-32.0)
Years with episode	1.0	(1.0-1.0)	6.0*	(2.0-17.0)	3.0*	(1.0-8.0)
12-month persistence	0.0		0.0	(0.0-0.0)	0.0	(0.0-1.0)
C. Total				, ,		` ′
Age-of-onset	18.0	(16.0-22.0)	18.0	(14.0-22.0)	21.0	(17.0-32.0)
Years with episode	1.0	(1.0-1.0)	5.0*	(2.0-15.0)	3.0*	(1.0-10.0)
12-month persistence	0.0		0.0	(0.0-1.0)	0.0	(0.0-1.0)
(n)	(23)	(52)	(1	15)

^{*} Significantly different from Anorexia Nervosa based on a .05 level, two-sided test.

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Table 4Difference in BMI categories at the time of interview in lifetime prevalence of DSM-IV disorders and related behavior

		Anorexia Nervosa	lervosa		Bulimia Ne	rvosa	Bir	ge-eating I	Disorder		reshold Bi	nge-eating	Any b	inge-eatin	g behavior
	%	OR	OR (95% CI)	%	OR	OR (95% CI)	%	OR	% OR (95% CI)		OR	% OR (95% CI) % OR (95% CI)	%	OR	(95% CI)
Current BMI	П														
	15.6	5.6	(0.9-33.4)	3.5	1.0	1.0 (0.2–5.3)	0.0			0.0			8.0	0.4	0.8 0.4 (0.1–1.7)
× 18.5	79.0	1.0		65.3	1.0		57.6	57.6 1.0**		72.1 1.0^^	\ \ \ \ \		63.8 1.0	1.0	
18.5-								2:			2:				
29.9 30-	5.3	0.3	0.3^{\wedge} (0.0–2.3)	20.9	1.0	1.0 (0.4–2.7)	27.6	1.7	27.6 1.7 (0.8–3.5)	24.4	1.1	24.4 1.1 (0.5–2.8)		1.3	24.8 1.3 (0.8–2.3)
39.9 > 40	0.0			10.3	2.6	(0.8–8.3)	14.8	*6.4	(2.2–11.0)	3.5	1.0	1.0 (0.3–3.9)	10.7	3.1*	(1.6–5.8)
χ^2_3 (p-		3.0	3.6(0.035)*		1.0	1.0 (0.412)		8.1	$8.1(0.001)^*$		0.0	0.0 (0.966)		4.7	4.7 (0.007)*

M BMI categories < 18.5 and 18.5–29.9 were collapsed

ABMI categories > 40 and 30–39.9 were collapsed

* Significant inter-cohort different based on a .05 level, two-sided test, controlling for age, sex and race-ethnicity.

Appendix table 4

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Cross-sectional socio-demographic profile of respondents with lifetime DSM-IV eating disorders and related behavior I

	Anoi OR	Anorexia Nervosa R (95% CI)	Bu	Bulimia Nervosa R (95% CI)	Binge-OR	Binge-eating Disorder OR (95% CI)	Subthre OR	Subthreshold Binge-eating OR (95% CI)	Any binge OR	Any binge-eating behavior OR (95% CI)
Race-ethnicity Non-Hispanic White	1.0	1	1.0	:	1.0	;	1.0	;	1.0	1
Non-Hispanic Black Hispanic	0.32	(0.1–1.4)	1.0	(0.4-2.5) $(0.4-11.0)$	0.7	(0.3–1.2)	1.9	(0.5-7.1) (0.2-5.2)	0.9	(0.4-1.7) $(0.6-2.1)$
Other		(0.1-3.9)		(0.8-4.5)	•	(0.1–2.5)	0.8	(0.2–3.2)		(0.3–1.8)
λ 3 (P-vaine) Education		(4677) (77		3.2 (.332)	•	(77+.) 0.7		(667.) 6.1	2	(600) 07
Less than high school	0.5	(0.1-2.6)	1.3	(0.4-4.1)	2.1	(0.8-5.2)	1.9	(0.4-10.0)	1.8	(1.0-3.4)
High school graduate	8.0	(0.4-1.7)	0.4	(0.2-0.9)	0.7	(0.3-1.8)	1.7	(0.5-6.7)	6.0	(0.5-1.6)
Some post-HS education	1:1	(0.4-3.3)	1.2	(0.6-2.7)	1.5	(0.8-3.1)	2.2	(0.7-7.0)	1.5	(0.9-2.4)
College graduate	1.0	:		١,	1.0	:	1.0	1		١,
χ^2_3 (p-value)		1.9 (.601)		8.4 (.039)*	T	5.5 (.002)		2.1 (.561)	17	17.4 (.001)*
Marital status										
Previously married	1.0	(0.2-4.1)	1.8	(0.8-3.9)	1.0	(0.5-1.8)	8.0	(0.3-1.7)	1.0	(0.7-1.6)
Never married	0.4	(0.1-2.0)	0.2	(0.1-0.5)	8.0	(0.6-1.3)	9.0	(0.3-1.2)	0.7	(0.5-0.9)
Married-cohabitating	1.0	1	1.0	1,	1.0	;	1.0	;		1
χ^2_2 (p-value)		1.8 (.411)		15.3 (.000)*	_	0.8 (.669)		2.1 (.347)	7.	7.2 (.027)*
Employment status										
Employed	1.0	1	1.0	;	1.0	;	1.0	1	1.0	1
Student	3.7	(0.4-37.0)	9.0	(0.1-5.3)	1.2	(0.3-5.5)	4.2	(0.7-24.6)	2.0	(0.9-4.3)
Homemaker	4.3	(1.4-12.7)	1.0	(0.4-2.2)	0.7	(0.3-1.9)	0.7	(0.1-3.2)	0.7	(0.3-1.3)
Retired			7.5	(0.4-142.1)	1.8	(0.2-14.8)	0.2	(0.0-0.7)	0.8	(0.2-3.6)
Other	$^{6.0}$	(0.1-6.3)	2.5	(1.1-5.8)	1.9	(0.9-3.8)	2.1	(0.6-7.5)	2.1	(1.4-3.2)
χ^2_4 (p-value)		13.9 (.003)		6.8 (.145)	71	5.5 (.240)		9.7 (.046)	15	(100.) 6.61

^{*} Significant at the .05 level, two-sided test

 $^{^{}I}$ Controlling for age and sex in every model

²Non-Hispanic Blacks and Hispanics were collapsed in this equation because of sparse data.

 $^{^3}$ Variables Retired and Other were collapsed in this equation because of sparse data.

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Table 5 Impairment in role functioning (Sheehan Disability Scales) associated with 12-month DSM-IV eating disorders and related behavior

	Bulimia Nervosa %	ervosa (se)	Binge-eating Disorder % (se)	g Disorder (se)	Subthreshold I	Binge-eating (se)	Any binge-ea	Any binge-eating behavior % (se)
I. Prevalence of any impairment	ent							
Home		(16.4)	46.1	(7.1)	18.4	(6.7)	38.1	(5.8)
Work	59.2	(14.8)	38.4	(6.8)	16.3	(8.5)	35.5	(6.1)
Personal life	47.2	(12.3)	39.7	(8.8)	15.5	(9.1)	34.3	(0.9)
Social life	54.1	(14.0)	58.8	(7.5)	18.9*	(6.7)	47.3	(6.2)
Any	78.0	(12.4)	62.6	(T.T)	21.8*	(10.3)	53.1	(9.9)
II. Prevalence of severe impa	irment							
Home	25.9	(15.5)	9.4	(4.2)	2.9	(3.0)	10.1	(3.9)
Work	25.9*	(15.5)	0.0	. 1	0.0	. 1	3.8	(2.8)
Personal life	9.9	(4.7)	5.7	(3.6)	1.6	(1.7)	4.7	(2.2)
Social life	31.7	(13.5)	15.9	(6.5)	1.6	(1.7)	14.3	(4.2)
Any	43.9	(16.3)	18.5	(6.8)	4.6	(3.4)	18.4	(5.3)
(u)	(16)		(51)	_	(21)	_	(88)	

* Significantly difference from Binge-eating Disorder based on a .05 level, two-sided test.

 Table 6

 Lifetime co-morbidity (OR) of DSM-IV Eating Disorders with other core NCS-R/DSM-IV disorders and related behaviors¹
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	 	Anorexia Nervos	ervosa	Bu	Bulimia Nervosa	.0Sa	Bin	Binge-eating Disorder	Disorder	Subth	reshold B	Subthreshold Binge-eating		Any binge-	
I. Anxiety disorders	%	OR	(95% CI)	%	OR	(95% CI)	%	OR	(95% CI)	%	OR	(95% CI)	3 %	ting behav OR	(95% CI)
Panic disorder	3.0	0.5	(0.1–2.7)	16.2	2.9*	(1.1)	13.2	2.9*	(1.8–4.8)	9.9	1.9	(0.7–5.4)	11.8	2.8*	(1.8–
Agoraphobia without panic	4.6	3.2	(0.6-	10.8	*6.8	(9.6) -0.6)	6.5	5.2*	(1.9–	6.0	*8.5	(1.5-	6.2	5.6	(2.4.5) (2.4.5)
Specific phobia	26.5	2.1*	(1.0-4.3)	50.1	5.4*	(2.6- (2.6-	37.1	3.7*	(2.26.5)	20.7	2.0	(0.9-4.3)	32.5	3.3*	(2.3– (2.3– (2.3–
Social phobia	24.8	2.1	(0.9–5.2)	41.3	*7.4	(2.7– (2.3)	31.9	3.2*	(2.1–4.9)	26.8	2.5*	(1.5-4.3)	31.0	3.2*	4.5) (4.5)
-	7.0	1.0	(0.3–3.5)	11.8	1.9	8.3) (0.9–	11.8	2.0*	(1.2–3.3)	6.6	2.3	(0.6–8.6)	11.9	2.2*	(5.1) -2.5.
Generalized anxiety disorder Post-	12.0	1.6	(0.5–5.7)	45.4	10.2*	(5.2– (5.2–	26.3	5.1*	(2.8–9.4)	5.6	1.1	(0.4–2.8)	20.2	*0.4	3.2) (2.6–
Obsessive-	0.0	÷	:	17.4	7.5*	(1.7– (1.7– 37.5)	8.2	*2.5	(1.4–	0.0	:	:	7.2	*9.7	(1.7–7.0
compuisive disorder Separation anxiety disorder	7.5	1.7	(0.4–6.3)	15.7	3.5*	(5.15) (5.45)	12.2	3.3*	(1.7-6.3)	2.6	8.0	(0.2–3.5)	6.7	2.7*	(1.6-
Any anxiety disorder ⁴	47.9	1.9	(0.9–4.1)	9.08	*9.8	9.0) (3.4– 21.6)	65.1	4.3*	(2.6–7.1)	40.4	2.0	(1.0-4.0)	59.5	3.7*	4.8) (2.5– 5.5)
II. Mood disorders Major depressive disorder	39.1	2.7*	(1.3–5.7)	50.1	*6.7	(1.7–	32.3	2.2*	(1.3–3.7)	17.7	1.3	(0.6–2.7)	30.9	2.2	(1.6 -
Dysthymia	12.8	*5.4	(1.2–	12.7	*4.4	(1.5– (1.5–	9.6	3.6*	(1.9–6.9)	5.5	2.9	(0.9-9.0)	9.1	3.8*	(2.2) -4.2)
Bipolar I-II disorders	3.0	8.0	(0.2-3.7)	17.7	*7.4	(2.1– (2.1–	12.5	3.6*	(2.1–6.3)	10.5	3.0	(8.6–6.0)	12.0	3.5*	6.1) (2.0–
Any mood disorder	42.1	2.4*	(1.2–4.7)	70.7	7.8*	(3.6-)	46.4	3.1*	(1.9–4.8)	28.2	1.8	(0.8–3.9)	44.0	3.0*	(2.1 ₋
III. Impulse-control disorders	4.7	1.0	(0.1–8.6)	3.6	0.7	(0.1–	9.4	2.2*	(1.0-4.6)	5.4	1.0	(0.2–4.1)	7.2	1.5	(0.8–
Intermittent explosive disorder Attention-deficit/	16.2	2.1	(0.5–9.7)	34.9	*4.8	5.6) (3.1–	19.8	3.1*	(1.5–6.2)	10.7	1.3	(0.5–3.8)	19.0	3.0*	3.1)
hyperactivity disorder Oppositional-	10.5	1.4	(0.2–8.1)	26.9	5.1*	(2.1– (2.1–	18.9	2.9	(1.4–6.3)	6.9	6.0	(0.2–3.5)	15.6	2.4	(1.3– (1.3–
defiant disorder ² Conduct disorder ²	8.6	1.2	(0.2–6.7)	26.5	*7.4	(2.0-	20.0	2.6	(1.4-4.8)	7.0	0.5	(0.1–2.2)	17.2	1.9	(1.0 c
Any impulse- control disorder	30.8	1.4	(0.4–5.3)	63.8	e.7*	(3.0– 15.2)	43.3	2.5*	(1.4–4.6)	22.3	8.0	(0.4–1.7)	40.2	2.2	3.5) 3.5)
Alcohol abuse or	24.5	2.9*	(1.2–7.2)	33.7	*9.7	(2.1–	21.4	2.2*	(1.2–4.0)	30.0	2.3	(1.0–5.6)	25.8	2.6	(1.9–
Alcohol dependence	12.3	3.0*	(1.3–7.0)	22.7	6.3	(2.3– 17.3)	12.4	2.7*	(1.1-6.7)	21.4	*1.7	4.1)	15.7	3.5*	(2.1)
Illicit drug abuse or	17.7	3.4*	(1.3–8.4)	26.0	5.3	(1.6- (1.6-	19.4	3.4*	(1.9–6.1)	22.9	2.9	(1.3–6.4)	20.2	3.4*	(2.2)
dependence Illicit drug dependence	5.2	2.2	- 4 .0)	15.0	*0.8	(1.5–	10.6	*6.4	(2.2–	13.5	*5.4	(1.5–	11.0	*6.4	(2:8) -8-2)
Any substance use disorder	27.0	3.0*	(1.2-7.1)	36.8	*9.7	(2.0– (2.0–	23.3	2.1*	(1.2-3.8)	35.5	2.8*	(1.2-6.5)	28.7	2.8*	(1.9) -6.1)
V. Any disorder						10.0)									0.5)

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	¥	Anorexia Nervosa	lervosa	Bū	Bulimia Nervosa	osa	Bin	Binge-eating Disorder	Disorder	Subth	reshold Bi	Subthreshold Binge-eating		Any binge-	
I. Anxiety disorders	%	% OR (95'	(95% CI)	%	OR	(95% CI)	%	OR	OR (95% CI)	%	% OR	(95% CI)	%	eating Denavior % OR (95% CI)	(95% CI)
Any disorder	56.2	56.2 1.3	(0.6–3.1)	94.5 17.6*	17.6*	(4.5-	78.9	78.9 4.2*	(2.2–7.9) 63.6 2.0	63.6	2.0	*(0.8–4.6) 76.5 3.7	76.5	3.7*	(2.3–
Exactly one	8.4	0.5	(0.1-1.8)	6.1	2.8	08.4) (0.7–	20.2	2.7*	(1.1–6.5)	9.4	0.7	(0.2–2.5)	15.9	1.9	(1.0) (1.0)
Exactly two	14.1	1.4	(0.5-4.5)	24.0	19.2	(4.2– (8.2–	8.6	2.2*	(1.0-4.6)	23.5	3.2*	(1.0-	15.7	3.2*	5.8) (1.6–
Three or more	33.8	2.3	(0.9–5.4)	64.4	33.7*	88.0) (8.7– 131.1)	48.9	*9.7	(4.0– 14.5)	30.7	2.6	(1.1-5.9)	45.0	6.4 *	6.3) (3.9– 10.5)

* Significant at the .05 level, two-sided test.

I A separate logistic regression equation was estimated for each comorbid disorder and a single equation was estimated for number of comorbid disorders. Each equation controlled for respondent age (in five-year intervals), sex, and race-ethnicity.

 $^2\mathrm{Restricted}$ to respondents in the age range 18–44 (n=1672)

3 Restricted to a random sub-sample of respondents (n=1139)

 $^{4}\mathrm{OCD}$ was coded as absent among respondents who were not assessed for this disorder.

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Age-of-onset priority of DSM-IV eating disorders and related behavior with comorbid DSM-IV disorders

					Percent w	where eati	ng disorder	ers or behav	ior are ten	nporally pr	imary ¹				
	An	Anorexia Nervo	70sa	Bu	limia Nervo	Sa	Binge-	eating Disc	order	Subthre	shold Binge	-eating	Any bin	nge-eating b	ehavior
	%	(se)	(n)	%	(se)	(n)	%	(se)	(n)	%	(se)	(n)	%	% (se)	(n)
Anxiety disorders	15.3	(11.3)	(14)	21.2	(8.9)	(42)	8.6	(3.3)	(9L)	4.0	(3.8)	(27)	13.0	(3.8)	(127)
Mood disorders	70.1	(14.8)	(14)	64.8	(9.1)	(36)	34.6	(9.1)	(28)	39.6	(13.5)	(20)	48.9	(5.9)	(100)
Impulse-	0.0	1	(9)	2.6	(2.2)	(24)	4.7	(3.2)	(45)	12.5	(7.6)	(15)	5.8	(2.3)	(75)
control disorders Substance use disorders	61.8	(16.9)	(6)	71.0	(12.1)	(18)	26.3	(8.2)	(31)	50.0	(14.7)	(17)	52.7	(8.2)	(62)

 $I_{
m Based}$ on comparison of retrospective AOO reports for eating disorders and the earliest comorbid disorder in the category

	Anoros	Anorovia Norvasa	Rulimio	Rulimia Normosa	Ringo-ootir	Bings_seting Disorder
	%	sa ivel vosa (se)	%	(se)	% %	(se)
$\frac{1}{1}$ I detime treatment l for any emotional problem	lem					
General medical		(12.6)	43.9	(11.4)	36.3	(5.6)
Psychiatrist	29.4	(12.8)	48.2	(8.1)	33.6	(6.3)
Other mental health	34.3	(10.2)	48.3	(6.6)	35.0	(3.2)
Human services	11.7	(7.5)	14.9	(3.0)	19.9	(4.2)
CAM	26.0	(12.1)	21.6	(6.1)	19.7	(4.1)
Any lifetime treatment	50.0	(14.6)	63.2	(7.8)	51.2	(6.5)
(u)		(23)	\$)	(52)		(115)
II. Twelve-month treatment I for any emotional problem 2						
General medical	•		26.1	(14.3)	20.8	(7.2)
Psychiatrist			0.6	(5.5)	5.0	(2.6)
Other mental health			3.7	(3.3)	24.0	(6.3)
Human services			13.5	(10.0)	7.1	(4.6)
CAM			0.0		12.4	(5.1)
Any 12-month treatment			15.6	(9.4)	28.5	(10.7)
(n)			(1)	(16)		(51)
III. Treatment of eating disorders						
Lifetime	33.8	(14.2)	43.2	(9.7)	43.6	(6.2)
Twelve-month			15.6	(9.4)	28.4	(6.4)

Igeneral medical treatment is treatment by a non-psychiatrist physician or nurse or other medical practitioner who is not a mental health specialist. Psychiatrist treatment is treatment by a psychiatrist. Other mental health treatment is treatment by any mental health specialist other than a psychiatrist (e.g., psychologist, psychiatric social worker). Human services treatment is treatment by a minister, priest, rabbi, or other spiritual advisor or by a caseworker in a social services agency. CAM (Complementary-alternative medical) treatment in a self-help group on treatment by an alternative medical provider (e.g., massage therapist, chiropractor).

 $[\]ensuremath{^{2}}$ There were no respondents with 12-month Anorexia Nervosa.

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 Table 8b

 Lifetime and 12-month treatment of DSM-IV eating disorders for females

		Anorexia Nervosa	Bulimia Nervosa	۱ ـ	Binge-eating Disorder	Disorder
	%	(se)	%	(se)	%	(se)
I. Lifetime treatment I for any emotional problem	blem					
General medical		(12.3)	49.4	(10.8)	42.2	(6.3)
Psychiatrist	30.0	(13.7)	40.8	(9.9)	33.1	(7.0)
Other mental health	24.7	(12.1)	56.4	(7.7)	38.4	(5.4)
Human services	14.6	(9.4)	16.3	(6.5)	21.3	(6.2)
CAM	20.0	(10.7)	24.8	(6.9)	24.5	(5.5)
Any lifetime treatment	37.7	(16.8)	59.8	(8.1)	52.3	(7.2)
(u)		(61)	(45)		(84)	
II. Twelve-month treatment for any emotional proble	2 m ²					
General medical	•		28.7	(15.3)	24.7	(7.8)
Psychiatrist			6.6	(5.9)	4.9	(2.8)
Other mental health			4.0	(3.5)	25.1	(10.2)
Human services			14.8	(10.9)	7.9	(6.2)
CAM			0.0		15.6	(6.8)
Any 12-month treatment			17.1	(10.2)	31.6	(10.7)
(n)			(15)		(39)	
III. Treatment of eating disorders						
Lifetime	29.8	(13.7)	47.0	(8.5)	50.8	(6.9)
I welve-month			1/.1	(10.2)	31.0	(10.7)

Igeneral medical treatment is treatment by a non-psychiatrist physician or nurse or other medical practitioner who is not a mental health specialist. Psychiatrist treatment is treatment by a psychiatrist. Other mental health treatment is treatment by any mental health specialist other than a psychiatrist (e.g., psychologist, psychiatric social worker). Human services treatment is treatment by a minister, priest, rabbi, or other spiritual advisor or by a caseworker in a social services agency. CAM (Complementary-alternative medical) treatment in a self-help group on treatment by an alternative medical provider (e.g., massage therapist, chiropractor).

 $\ensuremath{^{2}}$ There were no respondents with 12-month Anorexia Nervosa.

Lifetime and 12-month treatment of DSM-IV eating disorders for males

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		Anorexia Nervosa	Bulin	Bulimia Nervosa	Binge-eating Disorder	g Disorder
	%	(se)	%	(se)	%	(se)
I. Lifetime treatment ¹ for any emotional problem	blem					
General medical		(25.1)	22.9	(15.8)	24.3	(7.5)
Psychiatrist	26.9	(22.7)	75.8	(17.1)	34.8	(11.3)
Other mental health	73.1	(22.7)	18.0	(14.2)	28.1	(10.2)
Human services	0.0	1	9.6	(10.1)	17.1	(6.9)
CAM	50.2	(25.1)	9.6	(10.1)	10.1	(5.4)
Any lifetime treatment	100.0		75.8	(17.1)	48.8	(12.0)
(u)	•	(4)		(<i>L</i>)	(31)	
II. Twelve-month treatment for any emotional problem ²	nal problem ²					
General medical	•		0.0	1	12.2	(9.4)
Psychiatrist			0.0	1	5.4	(5.5)
Other mental health			0.0	1	21.5	(15.2)
Human services			0.0	1	5.4	(5.5)
CAM			0.0	1	5.4	(5.5)
Any 12-month treatment			0.0	I	21.5	(15.2)
(u)				(1)	(12)	
III. Treatment of eating disorders						
Lifetime	50.2	(25.1)	29.1	(19.3)	28.9	(9.4)
Twelve-month			0.0		21.5	(15.2)

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